



STIC Search Report

Biotech-Chem Library

STIC Database Tracking Number: 137610

TO: Ann Lam
Location: REM/3C70
Art Unit: 1641
Friday, November 12, 2004

Case Serial Number: 09/993314

From: Toby Port
Location: Biotech-Chem Library
REM-1A59
Phone: 571-272-2523
toby.port@uspto.gov

Search Notes

Examiner Lam,

Here are the results of the search you requested.

Please feel free to contact me if you have any questions.

Toby Port

Please note the structure as defined by the applicant did not achieve any relevant citations. I went on to do a dept search that found 7 citations, but they do not contain the structure described in the claim.



=> file reg; d stat que 111
FILE 'REGISTRY' ENTERED AT 12:58:46 ON 12 NOV 2004
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STRUCTURE FILE UPDATES: 10 NOV 2004 HIGHEST RN 778546-63-7
DICTIONARY FILE UPDATES: 10 NOV 2004 HIGHEST RN 778546-63-7

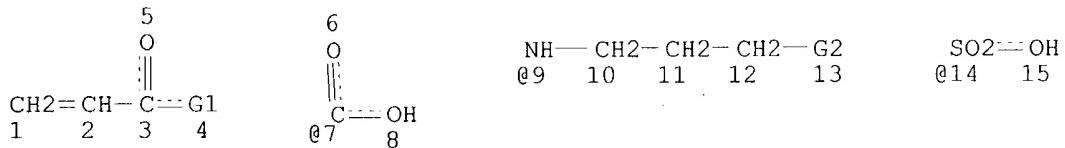
TSCA INFORMATION NOW CURRENT THROUGH MAY 21, 2004

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Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at:
<http://www.cas.org/ONLINE/DBSS/registryss.html>

L9 STR



Structures defined in claim 13

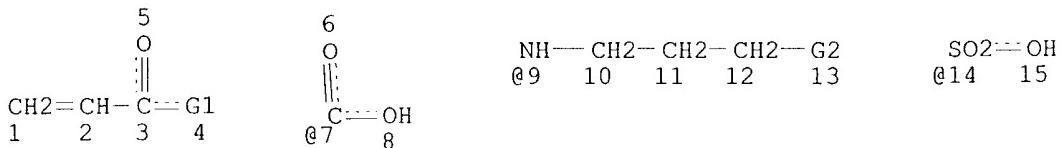
VAR G1=7/9
VAR G2=14/NH2
NODE ATTRIBUTES:
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
RING(S) ARE ISOLATED OR EMBEDDED
NUMBER OF NODES IS 15

STEREO ATTRIBUTES: NONE
L11 60 SEA FILE=REGISTRY SSS FUL L9

100.0% PROCESSED 304335 ITERATIONS 60 ANSWERS
SEARCH TIME: 00.00.02

=> d stat que 114
L9 STR



VAR G1=7/9

VAR G2=14/NH2

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 15

STEREO ATTRIBUTES: NONE

L11	60 SEA FILE=REGISTRY SSS FUL L9	↑
L13	5169 SEA FILE=REGISTRY ABB=ON PLU=ON PACR/PCT AND PA/PCT	
L14	0 SEA FILE=REGISTRY ABB=ON PLU=ON L11 AND L13	

*YL14 = Defined structures combined with
Polyacrylic (PACR) and Polyamide (PA)
in the PCT (Polymer Class Term) field*

=> file caplus; d que nos 116; d que nos 120; d que nos 121; d que nos 127
FILE 'CAPLUS' ENTERED AT 13:00:01 ON 12 NOV 2004

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FILE COVERS 1907 - 12 Nov 2004 VOL 141 ISS 20
FILE LAST UPDATED: 10 Nov 2004 (20041110/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

L9	STR
L11	60 SEA FILE=REGISTRY SSS FUL L9
L13	5169 SEA FILE=REGISTRY ABB=ON PLU=ON PACR/PCT AND PA/PCT
L15	9 SEA FILE=CAPLUS ABB=ON PLU=ON L11 AND L13
L16	0 SEA FILE=CAPLUS ABB=ON PLU=ON L15 AND MOBIL?

L9	STR
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L11 60 SEA FILE=REGISTRY SSS FUL L9
 L13 5169 SEA FILE=REGISTRY ABB=ON PLU=ON PACR/PCT AND PA/PCT
 L15 9 SEA FILE=CAPLUS ABB=ON PLU=ON L11 AND L13
 L20 0 SEA FILE=CAPLUS ABB=ON PLU=ON L15 AND ION EXCHANGE?

L9 STR
 L11 60 SEA FILE=REGISTRY SSS FUL L9
 L18 8424 SEA FILE=CAPLUS ABB=ON PLU=ON MOBILITY SHIFT ASSAY
 L21 0 SEA FILE=CAPLUS ABB=ON PLU=ON L11 AND L18

L9 STR
 L11 60 SEA FILE=REGISTRY SSS FUL L9
 L18 8424 SEA FILE=CAPLUS ABB=ON PLU=ON MOBILITY SHIFT ASSAY
 L27 0 SEA FILE=CAPLUS ABB=ON PLU=ON L11 AND L18

=> d que 119; d que 126
 L13 5169 SEA FILE=REGISTRY ABB=ON PLU=ON PACR/PCT AND PA/PCT
 L18 8424 SEA FILE=CAPLUS ABB=ON PLU=ON MOBILITY SHIFT ASSAY
 L19 2 SEA FILE=CAPLUS ABB=ON PLU=ON L18 AND L13

L13 5169 SEA FILE=REGISTRY ABB=ON PLU=ON PACR/PCT AND PA/PCT
 L22 128717 SEA FILE=CAPLUS ABB=ON PLU=ON ION EXCHANGE
 L25 286691 SEA FILE=CAPLUS ABB=ON PLU=ON MOBIL?
 L26 5 SEA FILE=CAPLUS ABB=ON PLU=ON L13 AND L22 AND L25

=> s 119 or 126
 L28 7 L19 OR L26

=> d ibib ab 128 1-7

L28 ANSWER 1 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 2004:62804 CAPLUS
 DOCUMENT NUMBER: 140:316145
 TITLE: Analysis of relative substances "in-1", "n-2" and
 "n-3" synthetic phosphorothioate oligonucleotides with
 IE-HPLC and PAGE
 AUTHOR(S): Li, Qilin; Zhou, Jian; Wang, Xiaoxing; Gao, Xiaoping
 CORPORATE SOURCE: Institute of Materia Medica, Chengdu Di Ao Group,
 Chengdu, 610041, Peop. Rep. China
 SOURCE: Yaowu Fenxi Zazhi (2002), 22(5), 371-375
 CODEN: YFZADL; ISSN: 0254-1793
 PUBLISHER: Yaowu Fenxi Zazhi Bianji Weiyuanhui
 DOCUMENT TYPE: Journal
 LANGUAGE: Chinese
 AB The relative substances for the synthesized phosphorothioate
 oligonucleotides of n=20-mer were analyzed using ion
 exchange HPLC and polyacrylamide gel electrophoresis (PAGE). The
 ion-exchange column was Gen-PakFAX (4.6 mm x 100 mm),
 The mobile phase A was 62.5 mmol.L-1 Tris.Cl, pH 8.15; the
 mobile phase B was 62.5 mmol.L-1 Tris.Cl, 2.5 mol.L-1 LiCl, pH
 8.15; the mobile phase C was 100% acetonitrile. The condition

of gradient elution was B: 30% → 50% 30 min, and C: 20%. The flow rate was 0.75 mL·min⁻¹, the detection was done at 260 nm. PAGE condition were 20% polyacrylamide and constant power at 25 W to electrophoresis. The relative substances n-1, n-2 and n-3 for the synthesized phosphorothioate oligonucleotides could be separated one by one by using the **ion-exchange** HPLC anal. method, which had reference value for the purification and the anal. of phosphorothioate oligonucleotides.

L28 ANSWER 2 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 2003:377085 CAPLUS
 DOCUMENT NUMBER: 138:380383
 TITLE: Methods and kits for detecting polymorphisms in nucleic acids using reverse phase HPLC or **ion-exchange** chromatography
 INVENTOR(S): Legendre, Benjamin, Jr.; Rudolph, Joseph G., III; Marino, Michael A.
 PATENT ASSIGNEE(S): Transgenomic, Inc., USA
 SOURCE: PCT Int. Appl., 62 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003040411	A1	20030515	WO 2002-US35409	20021104
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR				
US 2004035793	A1	20040226	US 2002-288406	20021104
EP 1451350	A1	20040901	EP 2002-778731	20021104
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK				
PRIORITY APPLN. INFO.:			US 2001-338627P	P 20011105
			US 2001-338041P	P 20011204
			US 2002-370749P	P 20020405
			WO 2002-US35409	W 20021104

AB Methods, systems, compns. and kits for improved detection of polynucleotides. In one aspect, there is provided a method for separating polynucleotides (such as DNA or RNA) using a liquid chromatog. separation device

(such as a reverse phase column or an **ion exchange** column), contacting eluted polynucleotides with intercalating dye, and detecting (such as by fluorescence detection) dye bound to the eluted polynucleotides. The invention preferably uses a post-column reactor, such as a mixing tee, downstream of the separation column. Sensitivity of mutation detection by denaturing high performance liquid chromatog. (DHPLC) is enhanced.

REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L28 ANSWER 3 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 2003:301189 CAPLUS

DOCUMENT NUMBER: 138:315802
 TITLE: Methods and kits for detection of nucleic acid polymorphisms using temperature compression denaturing high performance liquid chromatography
 INVENTOR(S): Taylor, Paul D.
 PATENT ASSIGNEE(S): Transgenomic, Inc., USA
 SOURCE: PCT Int. Appl., 75 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003031580	A2	20030417	WO 2002-US32042	20021007
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR				
US 2003225261	A1	20031204	US 2002-266906	20021007
PRIORITY APPLN. INFO.:			US 2001-327613P	P 20011005
			US 2001-335478P	P 20011101

OTHER SOURCE(S): MARPAT 138:315802
 AB Methods, compns., and kits for separating heteroduplex and homoduplex DNA mols. in a test mixture by temperature-compression denaturing high performance liquid chromatog. (tcDHPLC). The method includes use of nitrogen-containing additives in the **mobile** phase that allow detection of diverse heteroduplex mols. to be performed at the same pre-selected temperature. An example of a preferred additive is betaine. Standard mixts. of DNA fragments, such as mutation stds. containing known heteroduplex and homoduplex mols., can be used to select the concentration of additive and temperature. Compns. and kits including the **mobile** phase, mutation stds., PCR primers, separation media, and DNA polymerase are also provided.

L28 ANSWER 4 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 1997:202274 CAPLUS
 DOCUMENT NUMBER: 126:303376
 TITLE: Factors that affect the stability of protein-DNA complexes during gel electrophoresis
 AUTHOR(S): Fried, Michael G.; Bromberg, Jennifer L.
 CORPORATE SOURCE: Department of Biochemistry and Molecular Biology, The Pennsylvania State University College of Medicine, Hershey, PA, 17033, USA
 SOURCE: Electrophoresis (1997), 18(1), 6-11
 CODEN: ELCTDN; ISSN: 0173-0835
 PUBLISHER: VCH
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 AB The gel electrophoresis **mobility shift assay** is widely used for qual. and quant. characterization of protein complexes with nucleic acids. Often it is found that complexes persist within electrophoresis gels for much longer than expected on the basis of their free-solution lifetimes. Volume exclusion, direct interaction with gel

matrixes and the reduction of water activity by the gel have been proposed as mechanisms enhancing the stability of complexes during electrophoresis. We have used the well-characterized interaction of the *E. coli* cAMP receptor protein (CAP) with lactose promoter DNA to test these proposals. We found that the activity of water within polyacrylamide gels differs little from that of the buffer in which they were cast and that the dependence of the dissociation rate constant on water activity is too small for osmotic stabilization to contribute significantly to the lifetimes of CAP-DNA complexes. In addition, we found that a cross-linked gel matrix is not required for the stabilization of CAP-DNA complexes, that comparable stabilization is produced by three dissimilar polymers (linear polyacrylamide, dextran and polyethylene glycol), and that these polymers stabilize complexes more effectively than equivalent weight concns. of their cognate monomers. While these results challenge the notion that direct interaction with the gel matrix contributes to the stability of protein-DNA complexes, they are all features expected of excluded volume mechanisms.

L28 ANSWER 5 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1995:736175 CAPLUS
DOCUMENT NUMBER: 123:322990
TITLE: Effect of soluble aluminum ions on polyelectrolyte-alumina interaction. Kinetics of polymer adsorption and colloid stabilization
AUTHOR(S): Rignenbach, Eric; Chauvetea, Guy; Pefferkorn, Emile
CORPORATE SOURCE: Institut Charles Sadron, Strasbourg, 67083, Fr.
SOURCE: Colloids and Surfaces, A: Physicochemical and Engineering Aspects (1995), 99(2/3), 161-73
CODEN: CPEAEH; ISSN: 0927-7757
PUBLISHER: Elsevier
DOCUMENT TYPE: Journal
LANGUAGE: English
AB The authors studied the adsorption of high-mol.-weight polyacrylamide, hydrolyzed polyacrylamide, and polyacrylic acid on partially soluble colloidal Al₂O₃ in aqueous suspensions containing 10⁻³ N KCl and 3 × 10⁻⁴ N AlCl₃. The polymer solution and the colloidal suspension were mixed at pH 5.0, and the electrophoretic mobility and pH were recorded as a function of time. The polymer-Al ion interaction resulted in polymer ionization and complexation between carboxylic acid groups and Al ions characterized by a maximal degree of complexation close to 0.6. When the polymer was added to the oxide suspension, the authors also determined the concentration of the different species by potentiometric titration, and characterized the polyelectrolyte adsorption kinetics on colloidal Al₂O₃ by determining the variation with time of the amount of free and complexed polymers segments in the supernatant liquid phase. Two different situations were studied depending on the ratio of carboxylic acid to dissolved Al ions. For a low value of the ratio, the polymer was adsorbed quickly in a form which was highly complexed by Al ions pre-existing in the solution. For a high value of the ratio, the adsorption was very slow. Before adsorption, the polyacid also underwent an Al-H ion-exchange, the extent of which depended on the rate of oxide dissoln.; it appeared that adsorption increased very strongly with complexation. From an study of the colloidal stability of the system and the zeta-potential of the colloid-polymer complex as a function of the amount of polymer added to the solution, the domain of electrosteric stabilization was determined and the instability near the point of zero charge was attributed to the electrostatic attraction between the pos. and neg. charged groups.

L28 ANSWER 6 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 1995:483814 CAPLUS
 DOCUMENT NUMBER: 122:260341
 TITLE: Alternative to polyacrylamide gels improves the electrophoretic **mobility shift assay**
 AUTHOR(S): Vanek, P. G.; Fabian, S. J.; Fisher, C. L.; Chirikjian, J. G.; Collier, G. B.
 CORPORATE SOURCE: Georgetown Univ. Med. Cent., Washington, DC, USA
 SOURCE: BioTechniques (1995), 18(4), 704-6
 CODEN: BTNQDO; ISSN: 0736-6205
 PUBLISHER: Eaton
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 AB The authors outline a simplified protocol for the electrophoretic **mobility shift assay** utilizing TreviGel 500, a nontoxic alternative to polyacrylamide. The TreviGel 500 matrix combines the strength and resolution of polyacrylamide with the simplicity and flexibility of agarose in the casting of gels. Therefore, this method provides a simple, rapid and nontoxic alternative to current protocols for the investigation of protein:DNA interactions.

L28 ANSWER 7 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 1986:226207 CAPLUS
 DOCUMENT NUMBER: 104:226207
 TITLE: Grafting of poly(methacrylic acid) onto polycaproamide and the production of modified fibers with **ion-exchange** properties
 AUTHOR(S): Bogoeva-Gatseva, G.; Gabrielyan, G. A.; Gal'braikh, L. S.
 CORPORATE SOURCE: USSR
 SOURCE: Khimicheskie Volokna (1986), (2), 24-6
 CODEN: KVLKA4; ISSN: 0023-1118
 DOCUMENT TYPE: Journal
 LANGUAGE: Russian
 AB Methacrylic acid was graft polymerized on nylon 6 fibers in the presence of the catalytic system K₂S₂O₈-Na₂S₂O₃-Cu ion aquo or organic ligand complex. The system was most active when the fibers were treated with Cu complex prior to polymerization, and the optimal content of the complex was 0.001-0.002% (based on fiber weight). Organic complexes were more effective than aquo, and the phthalocyanine complex was the most effective among the former. Cu phenanthroline complex in combination with Na₂S₂O₃ and K₂S₂O₈ in 2.5:1 ratio fully inhibited the polymerization. The initial rate of grafting was higher for unoriented fibers, due to increased swelling, but the yield of graft copolymer was higher for oriented fibers, due to inhibited chain termination resulting from decreased chain **mobility**. The **ion-exchange** capacity of the modified fibers increased to 5.85 mequiv/g as graft degree rose to 54.6%.

=> => file caold; d que nos 129
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FILE COVERS 1907-1966

FILE LAST UPDATED: 01 May 1997 (19970501/UP)

This file contains CAS Registry Numbers for easy and accurate substance identification. Title keywords, authors, patent assignees, and patent information, e.g., patent numbers, are now searchable from 1907-1966. TIFF images of CA abstracts printed between 1907-1966 are available in the PAGE display formats.

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L29 0 SEA FILE=CAOLD ABB=ON PLU=ON L11

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=>